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## IN THE CLAIMS:

- 1. (Canceled)
- (Canceled)
- 3. (Canceled)
- (Withdrawn) The method of claim 1, wherein the at least one additional source includes a percent time in mode switch source.
- (Withdrawn) The method of claim 1, wherein the at least one additional source includes an R-wave and P-wave amplitude source.
- (Withdrawn) The method of claim 1, wherein the at least one additional source includes a reversion pace count source.
- (Withdrawn) The method of claim 1, wherein the at least one additional source includes a refractory sense count source.
- 8. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a high rate episode count source.
- 9. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a time from implant source.
- 10. (Canceled)
- 11. (Withdrawn) The method of claim 2, wherein the message indicates a lead conductor or connector issue.

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- 12. (Withdrawn) The method of claim 2, wherein the message indicates a lead insulation issue.
- 13. (Canceled)
- (Canceled)
- 15. (Withdrawn) The method of claim 13, wherein the biological interface issue includes lead dislodgement.
- (Withdrawn) The method of claim 13, wherein the biological interface issue includes exit block
- (Currently Amended) <u>A method of lead status monitoring in an</u> An
  implantable medical device (IMD) including a lead status monitoring system
  employing a method comprising the steps of:

collecting <u>lead impedance</u> data sets from a lead impedance source, collecting a stimulation threshold data source.

collecting data relating to one of a percent of time in mode switch, R-wave amplitude, P-wave amplitude, reversion pace count, refractory sense count, high rate episode count, and time from implant and at least one additional source included in the IMD: and

processing the <u>collected</u> data sets <u>in accordance with an algorithm having an integrated set of rules</u> to determine if a lead status event has occurred, wherein <u>each rule of the set applies a specific determination criterion to a particular aspect of the collected data</u> the at least one additional source includes one of a percent time in mode switch source, an R-wave and P-wave amplitude source, a reversion pace count source, a refractory sense count source, a high rate episode count source, and a time from implant source.

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- 18. (Previously Presented) The method of claim 17, further comprising providing a message indicating a lead-related condition to a user based on the lead status event.
- 19. (Previously Presented) The method of claim 18, wherein the message indicates one of a lead conductor or connector issue, a lead insulation issue, and a biological interface issue.
- (Previously Presented) The method of claim 19, wherein the biological interface issue includes one of myocardial perforation, lead dislodgement, and exit block.
- 21. (Previously Presented) The method of claim 17, wherein the processing comprises: assigning weighted values to the collected data sets; and summing the assigned weighted values to determine if one of a plurality of lead status events has occurred.